CLEAR EYES MAXIMUM REDNESS RELIEF- naphazoline hydrochloride and glycerin liquid Prestige Brands Holdings, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Clear Eyes Maximum Redness Relief

Drug Facts

Active ingredients

Glycerin 0.5%

Purpose

Lubricant

Active ingredients

Naphazoline hydrochloride 0.03%

Purpose

Redness Reliever

Uses

- for the relief of redness of the eye due to minor eye irritations.
- for the temporary relief of burning and irritation due to the dryness of the eye.
- for the use as a protectant against further irritation or dryness of the eye.

Warnings

For external use only.

Do not use

if solution changes color or becomes cloudy.

Ask a doctor before use if

you have narrow angle glaucoma.

When using this product:

- To avoid contamination, do not touch tip of container to any surface.
- Replace cap after using.
- Overuse may produce increased redness of the eye.
- Pupils may become enlarged temporarily.

Stop use & ask a doctor if:

- you experience eye pain
- you experience changes in vision

- you experience continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Instill 1 to 2 drops in the affected eye(s) up to four times daily

Other information

- Store at room temperature.
- Remove contact lenses before using.
- **Tamper Evident**: Do not use if neckband on bottle is broken or missing.

Inactive ingredients

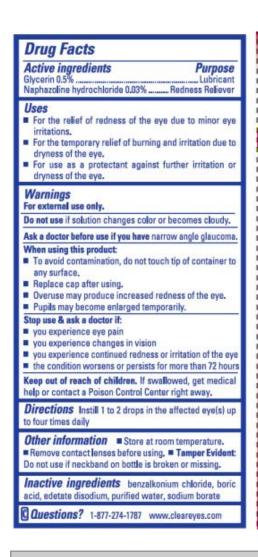
benzalkonium chloride, boric acid, edetate disodium, purified water, sodium borate

Questions?

1-877-274-1787 www.cleareyes.com

PRINCIPAL DISPLAY PANEL

Clear
eyes®
MAXIMUM
REDNESS RELIEF
LUBRICANT/REDNESS
RELIEVER EYE DROPS
Sterile 1.0 FL OZ (30 mL)





CLEAR EYES MAXIMUM REDNESS RELIEF

naphazoline hydrochloride and glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67172-494
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NAPHAZOLINE HYDRO CHLO RIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	.00015 mg in 1 mL	
GLYCERIN (UNII: PDC6 A3C0 OX) (GLYCERIN - UNII: PDC6 A3C0 OX)	GLYCERIN	.0025 mL in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
BORIC ACID (UNII: R57ZHV85D4)			
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)			
WATER (UNII: 059QF0KO0R)			

SODIUM BORATE (UNII: 91MBZ8H3QO)

Product Characteristics				
Color	WHITE	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-494-01	1 in 1 BOX		
1		30 mL in 1 BOTTLE, DROPPER		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part349	09/01/2010	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Establishment				
Name	Address	ID/FEI	Business Operations	
Altaire Pharmaceuticals Inc.		786790378	MANUFACTURE	

Revised: 9/2010 Prestige Brands Holdings, Inc.